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AMENDMENTS TO THE CLAIMS

1. (Original) A compound of formula (I) or a pharmaceutically acceptable derivative thereof

in which:

R¹ is bromo; and

 R^2 is halogen, C_{1-6} alkyl or C_{1-6} alkoxy.

- 2. (Original) The compound according to claim 1 in which R^2 is halogen or C_{1-6} alkoxy.
- 3. (Original) The compound according to claim 2 in which R² is fluoro, methoxy or ethoxy.

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4. (Currently Amended) The compound according to claim 1 which is selected from the group consisting of

- (S) 2-{[1-(2-Bromo-5-methylphenyl)methanoyl]amino} 3 (4'-cyano-2',6'-dimethoxybiphenyl-4-yl)propionic acid);
- (S)-2-{[(2-Bromo 5-chlorophenyl)methanoyl]amino}-3-[4' cyano-2',6'-dimethoxy)biphenyl-4-yl]propionic acid;
- (S) 2 {[(2,5-Dibromophenyl)methanoyl]amino} -3 [4' cyano 2',6' dimethoxy)biphenyl -4 yl]propionic acid;
- (S) 2-{[(5 (iso Propoxy) 2 bromophenyl)methanoyl]amino} 3 [4' cyano 2',6' dimethoxy)biphenyl-4-yl]propionic acid or a pharmaceutically acceptable derivative thereof.

The compound according to claim 1 which is selected from the group consisting of (S)-2-{[1-(2-Bromo-5-methylphenyl)methanoyl]amino}-3-(4'-cyano-2',6'-dimethoxybiphenyl-4-yl)propionic acid);

- (S)-2-{[(2-Bromo-5-chlorophenyl)methanoyl]amino}-3-[4'-cyano-2',6'-dimethoxybiphenyl-4-yl]propionic acid;
- (S)-2-{[(2,5-Dibromophenyl)methanoyl]amino}-3-[4'-cyano-2',6'-dimethoxybiphenyl-4-yl]propionic acid;
- (S)-2-{[(5-(iso-Propoxy)-2-bromophenyl)methanoyl]amino}-3-[4'-cyano-2',6'-dimethoxybiphenyl-4-yl]propionic acid or a pharmaceutically acceptable derivative thereof.

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5. (Orignal) (S)-2-{[1-(2-Bromo-5-ethoxyphenyl)methanoyl]amino}-3-(4'-cyano-2',6'-dimethoxybiphenyl-4-yl)propionic acid or a pharmaceutically acceptable derivative thereof.

- 6. (Original) (S)-2-{[1-(2-Bromo-5-fluorophenyl)methanoyl]amino}-3-(4'-cyano-2',6'-dimethoxybiphenyl-4-yl)propionic acid or a pharmaceutically acceptable derivative thereof.
- 7. (Original) (S)-2-{[1-(2-Bromo-5-methoxyphenyl)methanoyl]amino}-3-(4'-cyano-2',6'-dimethoxybiphenyl-4-yl)propionic acid or a pharmaceutically acceptable derivative thereof.

 or a pharmaceutically acceptable derivative thereof.
- 8. (Original) A process for the preparation of a compound of formula (I) which comprises hydrolyzing of a carboxylic acid ester derivative of formula (II):

in which R¹ and R² are as defined in formula (I) and R is a group capable of forming a carboxylic acid ester and optionally thereafter forming a pharmaceutically acceptable derivative thereof.

- 9. (Currently Amended) A compound according to any one of claims 1 to 7 claim 1 for use in therapy.
- 10. (Currently Amended) A pharmaceutical composition which comprises a therapeutically effective amount of a compound according to any one of claims 1 to 7 claim 1 in admixture with a pharmaceutically acceptable carrier or diluent.
- 11. (Currently Amended) A pharmaceutical composition comprising a compound according to any one of claims 1 to 7 claim 1 together with another therapeutically active agent.
- 12. (Currently Amended) A use of a compound according to any one of claims 1 to 7 claim 1 in the manufacture of a medicament for the treatment or prevention of conditions in which an inhibitor of α_4 integrin mediated cell adhesion is beneficial.
- 13. (Currently Amended) A method for the treatment or prevention of conditions in which an inhibitor of α_4 integrin mediated cell adhesion is beneficial which comprises administering to a patient in need thereof a safe and effective amount of a compound according to any one of claims 1 to 7 claim 1.

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(Original) The method according to claim 13, wherein said condition is selected from the 14. group consisting of rheumatoid arthritis (RA); asthma; allergic conditions such as rhinitis; adult respiratory distress syndrome; AIDS-dementia; Alzheimer's disease; cardiovascular diseases; thrombosis or harmful platelet aggregation; reocclusion following thrombolysis; reperfusion injury; skin inflammatory diseases such as psoriasis, eczema, contact dermatitis and atopic dermatitis; diabetes (e.g., insulin-dependent diabetes mellitus, autoimmune diabetes); multiple sclerosis; systemic lupus erythematosus (SLE); inflammatory bowel disease such as ulcerative colitis, Crohn's disease (regional enteritis) and pouchitis (for example, resulting after proctocolectomy and ileoanal anastomosis); diseases associated with leukocyte infiltration to the gastrointestinal tract such as Celiac disease, nontropical Sprue, enteropathy associated with seronegative arthropathies, lymphocytic or collagenous colitis, and eosinophilic gastroenteritis; diseases associated with leukocyte infiltration to other epithelial lined tissues, such as skin, urinary tract, respiratory airway, and joint synovium; pancreatitis; mastitis (mammary gland); hepatitis; cholecystitis; cholangitis or pericholangitis (bile duct and surrounding tissue of the liver); bronchitis; sinusitis; inflammatory diseases of the lung which result in interstitial fibrosis, such as hypersensitivity pneumonitis; collagen disease (in SLE and RA); sarcoidosis; osteoporosis; osteoarthritis; atherosclerosis; neoplastic diseases including metastasis of neoplastic or cancerous growth; wound (wound healing enhancement); certain eye diseases such as retinal detachment, allergic conjunctivitis and autoimmune uveitis; Sjogren's syndrome; rejection (chronic and acute) after organ transplantation; host vs. graft or graft vs. host diseases; intimal hyperplasia; arteriosclerosis (including graft arteriosclerosis after transplantation);

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reinfarction or restenosis after surgery such as percutaneous transluminal coronary angioplasty (PTCA) and percutaneous transluminal artery recanalization; nephritis; tumor angiogenesis; malignant tumor; multiple myeloma and myeloma-induced bone resorption; sepsis; and central nervous system injury such as stroke, traumatic brain injury and spinal cord injury and Meniere's disease.

- 15. (Original) The method according to claim 14, wherein said condition is inflammatory bowel disease or multiple sclerosis.
- 16. (Original) A compound of formula (II):

in which R¹ and R² are as defined in formula (I) and R is a group capable of forming a carboxylic acid ester.

- 17. (Original) A compound according to claim 16 in which R² is C₁₋₆alkoxy or fluoro.
- 18. (Original) A compound of formula (III) or an acid addition salt thereof:

in which R is a group capable of forming a carboxylic acid ester.

19. (Original) A compound of formula (VI):

(VI)